

Supplemental Material to:

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**B-Myb promotes S-phase independently of
its sequence-specific DNA binding activity and interacts
with polymerase delta interacting protein 1 (Pdip1)**

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Supplementary Materials

Supplementary Fig. 1 – B-Myb knock-down in HepG2 cells does not cause a G2/M-block. HepG2 cells were transfected with control or B-Myb specific siRNAs. 48 hours after transfection the cells were analyzed by flow cytometry.

Supplementary Fig. 2 – c-Myc expression is not affected by B-Myb knock-down. HepG2 cells were transfected with control or B-Myb specific siRNAs. 48 hours after transfection the cells were analyzed by western blotting with antibodies against B-Myb, c-Myc and β -actin.

Supplementary Fig. 3 – Pdip1 does not affect the transactivation potential of B-Myb. QT6 cells were transfected with expression vectors for full-length B-Myb, c-terminally truncated B-Myb (B-Myb Δ Sac) and Myc-Pdip1, as shown at the bottom. Additionally, cells were transfected with the Myb-responsive luciferase reporter gene 3xAtkluc and the β -galactosidase plasmid pCMV β . Cells were harvested 16 hours after transfection and analyzed for luciferase and β -galactosidase activity. The columns show the average luciferase activity normalized to the β -galactosidase activity. Thin lines show standard deviations.





